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**Listing of Claims:** 

The following listing of claims replaces all prior versions and listings of claims in the application:

- 1.-87. (Canceled)
- 88. (New) A polypeptide having the sequence SEQ ID NO:56.
- 89. (New) The polypeptide of claim 88, which is glycosylated.
- 90. (New) The polypeptide of claim 89, further comprising at least one PEG molecule covalently attached to the polypeptide.
- 91. (New) The polypeptide of claim 90, comprising one PEG molecule covalently attached to the polypeptide.
- 92. (New) The polypeptide of claim 91, wherein the PEG molecule has a molecular weight of about 12 kDa.
- 93. (New) The polypeptide of claim 91, wherein the PEG molecule has a molecular weight of about 20 kDa.
- 94. (New) A composition comprising the polypeptide of claim 89 and a pharmaceutically acceptable diluent, carrier, or excipient.
- 95. (New) A composition comprising the polypeptide of claim 93 and a pharmaceutically acceptable diluent, carrier, or excipient.

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96. (New) A nucleic acid comprising a nucleotide sequence encoding the polypeptide of claim 88.

- 97. (New) An expression vector comprising the nucleic acid of claim 96.
- 98. (New) A glycosylating host cell comprising the expression vector of claim 97.
- 99. (New) The glycosylating host cell of claim 98, wherein the host cell is a CHO cell.
- 100. (New) A method of making a polypeptide, the method comprising: providing a culture comprising a glycosylating host cell, the glycosylating host cell comprising a nucleotide sequence which encodes the polypeptide of claim 88, culturing the culture under conditions which permit expression and glycosylation of the polypeptide, and recovering the polypeptide.
- 101. (New) The method of claim 100, wherein the glycosylating host cell is a CHO cell.
- 102. (New) The method of claim 100, further comprising attaching at least one PEG molecule to the polypeptide.
- 103. (New) A method of treating a mammal with a disease for which interferon  $\beta$  is a useful treatment, the method comprising administering to the mammal an effective amount of the composition of claim 95.
- 104. (New) The method of claim 103, wherein the disease is multiple sclerosis.
- 105. (New) A polypeptide having the sequence SEQ ID NO:57.
- 106. (New) The polypeptide of claim 105, which is glycosylated.

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107. (New) The polypeptide of claim 106, further comprising at least one PEG molecule covalently attached to the polypeptide.

- 108. (New) The polypeptide of claim 107, comprising one PEG molecule covalently attached to the polypeptide.
- 109. (New) The polypeptide of claim 108, wherein the PEG molecule has a molecular weight of about 12 kDa.
- 110. (New) The polypeptide of claim 108, wherein the PEG molecule has a molecular weight of about 20 kDa.
- 111. (New) A composition comprising the polypeptide of claim 106 and a pharmaceutically acceptable diluent, carrier, or excipient.
- 112. (New) A composition comprising the polypeptide of claim 110 and a pharmaceutically acceptable diluent, carrier, or excipient.
- 113. (New) A nucleic acid comprising a nucleotide sequence encoding the polypeptide of claim 105.
- 114. (New) An expression vector comprising the nucleic acid of claim 113.
- 115. (New) A glycosylating host cell comprising the expression vector of claim 114.
- 116. (New) The glycosylating host cell of claim 115, wherein the host cell is a CHO cell.

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117. (New) A method of making a polypeptide, the method comprising: providing a culture comprising a glycosylating host cell, the glycosylating host cell comprising a nucleotide sequence which encodes the polypeptide of claim 105, culturing the culture under conditions which permit expression and glycosylation of the polypeptide, and recovering the polypeptide.

- 118. (New) The method of claim 117, wherein the glycosylating host cell is a CHO cell.
- 119. (New) The method of claim 117, further comprising attaching at least one PEG molecule to the polypeptide.
- 120. (New) A method of treating a mammal with a disease for which interferon  $\beta$  is a useful treatment, the method comprising administering to the mammal an effective amount of the composition of claim 112.
- 121. (New) The method of claim 120, wherein the disease is multiple sclerosis.